

**COLD AND FLU DAYTIME, NIGHTTIME, MULTI-SYMPTOM- acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl
Walgreen Company**

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Walgreens 44-659660-22-Delisted

Active ingredients (in each liquid-filled capsule) (Daytime Cold & Flu)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer
Cough suppressant
Nasal decongestant

Active ingredients (in each liquid-filled capsule) (Nighttime Cold & Flu)

Acetaminophen 325 mg
Dextromethorphan HBr 15 mg
Doxylamine succinate 6.25 mg

Purpose

Pain reliever/fever reducer
Cough suppressant
Antihistamine

Uses

- temporarily relieves common cold and flu symptoms:
 - fever
 - headache
 - sore throat
 - minor aches and pains
 - cough due to minor throat and bronchial irritation
 - nasal congestion (***Daytime only***)
 - runny nose and sneezing (***Nighttime only***)

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.
- if you have ever had an allergic reaction to this product or any of its ingredients
- to make a child sleepy (**Nighttime only**)

Ask a doctor before use if you have

- cough that occurs with too much phlegm (mucus)
- liver disease
- difficulty in urination due to enlargement of the prostate gland
- diabetes (**Daytime only**)
- heart disease (**Daytime only**)
- thyroid disease (**Daytime only**)
- high blood pressure (**Daytime only**)
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema (**Daytime only**)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis, or emphysema (**Nighttime only**)
- glaucoma (**Nighttime only**)

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers (**Nighttime only**)

When using this product

- **do not exceed recommended dosage**
- marked drowsiness may occur (**Nighttime only**)

- avoid alcoholic beverages (**Nighttime only**)
- excitability may occur, especially in children (**Nighttime only**)
- use caution when driving a motor vehicle or operating machinery (**Nighttime only**)
- alcohol, sedatives, and tranquilizers may increase drowsiness (**Nighttime only**)

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur (**Daytime only**)
- pain, nasal congestion, or cough gets worse or lasts more than 7 days (**Daytime only**)
- pain or cough gets worse or lasts more than 7 days (**Nighttime only**)
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts. These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Read each section carefully. Do not take DAYTIME and NIGHTTIME products at the same time.

Directions (Daytime only)

- **do not take more than directed**
- do not take more than 8 capsules per 24 hours
- adults and children 12 years and over: take 2 capsules with water every 4 hours
- children under 12 years: ask a doctor

Directions (Nighttime only)

- **do not take more than directed**
- do not take more than 8 capsules per 24 hours
- adults and children 12 years and over: take 2 capsules with water every 6 hours
- children under 12 years: ask a doctor

Do not take more than 8 capsules of Daytime and Nighttime products in any 24-hour period.

Other information

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F)

- protect from heat, humidity and light
- see end flap for expiration date and lot number

Inactive ingredients (Daytime only)

edible white ink, FD&C red #40, FD&C yellow #6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol

Inactive ingredients (Nighttime only)

D&C yellow #10, edible white ink, FD&C blue #1, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol

Questions or comments?

1-800-426-9391

Principal Display Panel

DAY & NIGHT PACK

NDC 0363-6596-22

Walgreens

Compare to Vicks® DayQuil® & NyQuil® Cold & Flu LiquiCaps® active ingredients††

DAYTIME • NON DROWSY Cold & Flu ACETAMINOPHEN / PAIN RELIEVER / FEVER REDUCER DEXTROMETHORPHAN HBr / COUGH SUPPRESSANT PHENYLEPHRINE HCl / NASAL DECONGESTANT MULTI-SYMPTOM ACTUAL SIZE 32 LIQUID CAPS	NIGHTTIME Cold & Flu ACETAMINOPHEN / PAIN RELIEVER / FEVER REDUCER DEXTROMETHORPHAN HBr / COUGH SUPPRESSANT DOXYLAMINE SUCCINATE / ANTIHISTAMINE MULTI-SYMPTOM ACTUAL SIZE 16 LIQUID CAPS
---	--

48 TOTAL LIQUID CAPS

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

Walgreens Pharmacist Recommended†

†Walgreens Pharmacist Survey

††This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademark Vicks® DayQuil® & NyQuil® Cold & Flu LiquiCaps®.
50844 REV0619B65966022

DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015

Walgreens

100% SATISFACTION GUARANTEED

walgreens.com ©2017 Walgreen Co.

PRODUCT OF CHINA
PACKAGED AND QUALITY
ASSURED IN THE U.S.A.

Walgreens 44-659660

COLD AND FLU DAYTIME, NIGHTTIME, MULTI-SYMPTOM

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-6596
--------------	----------------	--------------------	---------------

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-6596-22	1 in 1 PACKAGE, COMBINATION; Type 0: Not a Combination Product	03/01/2015	10/24/2022

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	4 BLISTER PACK	32
Part 2	2 BLISTER PACK	16

Part 1 of 2

COLD AND FLU DAYTIME, MULTI-SYMPTOM

acetaminophen, dextromethorphan hbr, phenylephrine hcl capsule, liquid filled

Product Information

Route of Administration	ORAL
-------------------------	------

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

SORBITOL (UNII: 506T60A25R)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	orange (clear)	Score	no score
Shape	OVAL	Size	21mm
Flavor		Imprint Code	659
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	03/01/2015	

Part 2 of 2

COLD AND FLU NIGHTTIME, MULTI-SYMPOM

acetaminophen, dextromethorphan hbr, doxylamine succinate capsule, liquid filled

Product Information

Route of Administration	ORAL
--------------------------------	------

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg

Inactive Ingredients

Ingredient Name	Strength
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics

Color	green (clear)	Score	no score
Shape	OVAL	Size	21mm
Flavor		Imprint Code	660
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	03/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	03/01/2015	10/24/2022

Labeler - Walgreen Company (008965063)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(0363-6596)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(0363-6596)

Establishment

Name	Address	ID/FEI	Business Operations
------	---------	--------	---------------------

LNK International, Inc.		832867837	pack(0363-6596)
-------------------------	--	-----------	-----------------

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	pack(0363-6596)

Revised: 11/2021

Walgreen Company